

AMENDMENTS TO THE CLAIMS

Please amend the claims as follows.

1. (Currently Amended) A pharmaceutical composition, comprising: ~~a safe and effective amount of (a)~~ at least one selected from the group consisting of the human liver regeneration associated protein hLRTM4, a polynucleotide comprising the hLRTM4 gene, and a polynucleotide comprising a degenerate sequence of the hLRTM4 gene; and (b) a pharmaceutically acceptable vehicle, diluent or carrier.
2. (Currently Amended) The composition of claim 1, wherein the hLRTM4 protein has the amino acid sequence as shown in SEQ ID NO:2, and the hLRTM4 gene has a sequence as shown in SEQ ID NO: 1.
3. (Currently Amended) The ~~composition~~ method of claim 4 ~~[[1]]~~, wherein the ~~safe and effective amount of hLRTM4~~ is 1 µg-5 mg/kg body weight per day.
4. (Currently Amended) ~~A use of liver regeneration associated protein hLRTM4 in the preparation of a drug used~~ A method for treating liver injury, comprising administering an effective amount of the pharmaceutical composition of claim 1 to a subject in need thereof.
5. (Currently Amended) The ~~use~~ method of claim 4, wherein the ~~drug is used to treat liver injury~~ is acute or chronic hepatitis, liver cirrhosis, or liver pathological changes caused by liver cancer.
6. (Currently Amended) A pharmaceutical composition, comprising: ~~a safe and effective amount of (a)~~ an antagonist[[s]] of hLRTM4 protein, hLRTM4 gene, or hLRTM4 gene transcript, wherein the hLRTM4 protein has a sequence of SEQ ID NO: 2 and the hLRTM4 gene has a sequence of SEQ ID NO: 1; and (b) a pharmaceutically acceptable vehicle, diluent or carrier.

~~wherein the antagonists are selected from the group consisting of: (i) an antisense polynucleotide to hLRTM4, wherein the polynucleotide has the antisense nucleotide sequence as shown in SEQ ID NO: 1 and has a length of 15-625 bp, (ii) small interfering double-strand RNAs of hLRTM4, wherein the RNAs have the nucleotide sequence as shown in SEQ ID NO: 1 and have a length of 17-23 bp and a 3' terminal dtdt sequence; and/or (iii) a specific antibody against hLRTM4, as well as a pharmaceutically acceptable vehicle, diluent or carrier.~~

7. (Currently Amended) ~~The composition of claim 6, wherein the polynucleotides has the full-length antisense sequence to SEQ ID NO: 1~~ the antagonist is at least one selected from the group consisting of: (i) an antisense polynucleotide for the hLRTM4 gene transcript, (ii) a small interfering RNA for the hLRTM4 gene transcript; and (iii) an antibody against the hLRTM4 protein.
8. (Currently Amended) ~~The composition~~ method of claim [[6]] 9, wherein the safe and effective amount of antagonist to hLRTM4 is 1 ug-5 mg/kg body weight per day.
9. (Currently Amended) ~~A use method for treating hepatocellular carcinoma or gastric adenocarcinoma, comprising administering to a subject in need thereof an effective amount of the composition of claim 6. of hLRTM4 protein antagonist for the preparation of a drug for treating hepatocellular carcinoma, wherein the antagonist is selected from: (i) an antisense polynucleotide to hLRTM4, wherein the polynucleotide has the nucleotide sequence as shown in SEQ ID NO: 1 and has a length of 15-625 bp; (ii) a small interfering double-strand RNA of hLRTM4, wherein the RNA has the nucleotide sequence as shown in SEQ ID NO: 1 and has a length of 17-23 bp and a 3' terminal dtdt sequence; and/or (iii) a specific antibody against hLRTM4.~~
10. (Currently Amended) ~~The use composition of claim 7 [[9]], wherein the antagonist is [[an]] the antisense polynucleotide having a length of 15 – 625 nucleotides. to hLRTM4, wherein~~

~~the polynucleotide has the nucleotide sequence as shown in SEQ ID NO: 1 and has a length of 15-625 bp.~~

11. (New) The composition of claim 7, wherein the antagonist is the small interfering RNA having a length of 16-23 bp and a 3' terminal dTdT sequence.
12. (New) An isolated hLRTM4 protein or a fragment thereof, wherein the hLRTM4 protein has a sequence of SEQ ID NO: 2.
13. (New) An isolated polynucleotide encoding the protein or the fragment of claim 12.
14. (New) The isolated polynucleotide of claim 13, wherein the isolated polynucleotide has a sequence of SEQ ID NO: 1.
15. (New) The isolated polynucleotide of claim 13, wherein the isolated polynucleotide is incorporated in an expression vector.